

EXHIBIT 1

From: Misbah Sherwani <MSherwani@actavis.com>
Sent: Tuesday, April 15, 2008 2:50 PM
To:
Subject: Fw: List by Product
Attach: 5-Sept 07 - Present Investigations by product.xls

She's gone through both.....there should be no more surprises.

----- Forwarded by Misbah Sherwani/QA/ELZ/Alpharma on 04/15/2008 02:48 PM -----

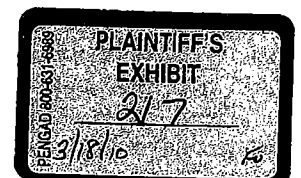
Michael Ponzo/Actavis@Exchange
04/15/2008 02:17 PM

To
Misbah Sherwani/QA/ELZ/Alpharma@Alpharma
cc
Anthony Castellazzo/Actavis@Exchange
Subject
List by Product

There were 2 that were incorrect (did not state stability OOS). Investigations
07-059 & 07-101.
All stability OOS are now highlighted.

Mike

<<...>>

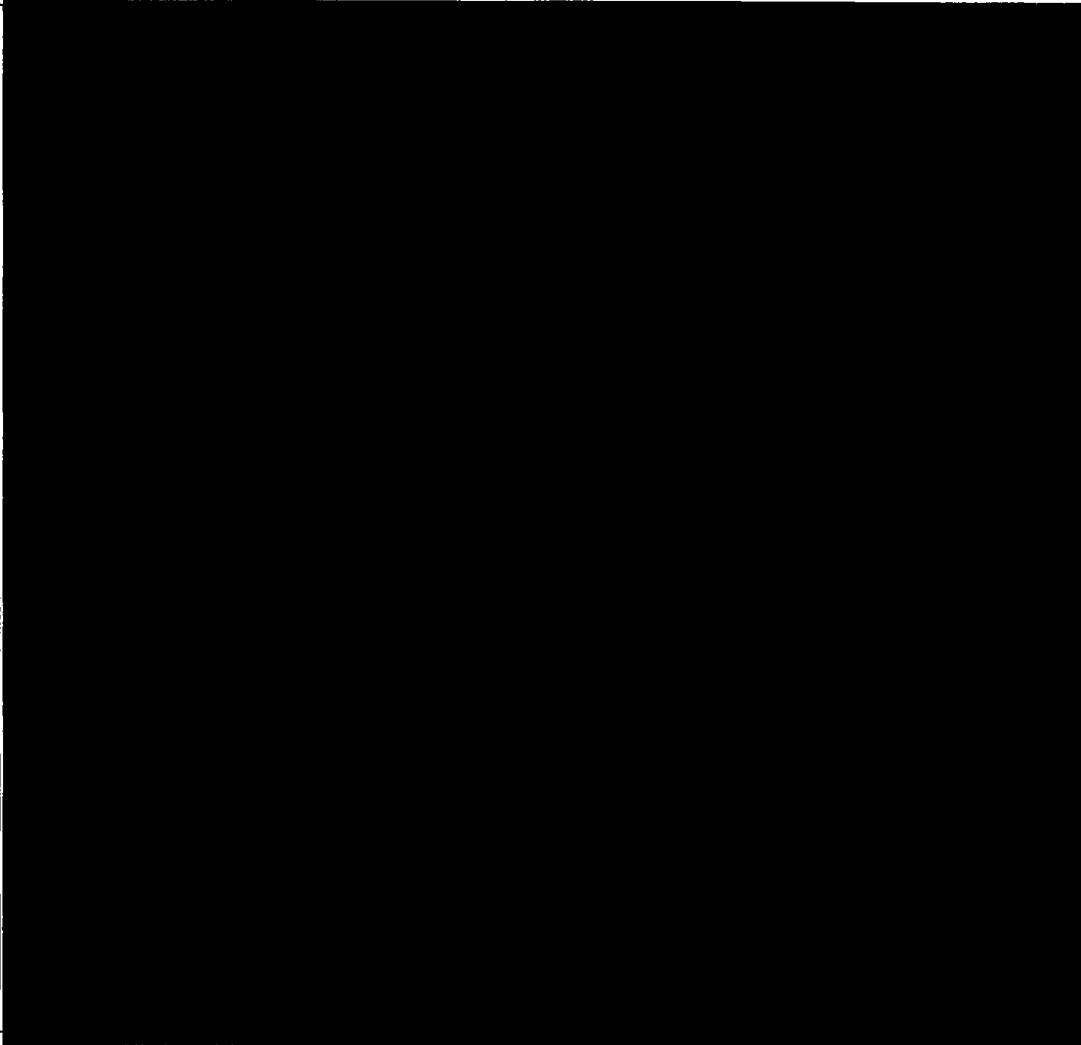


Investigation Number	Date Issued	Lot Number	Product No.	Product Name	Reason for Investigation	Status	Closed Date
[REDACTED]							

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[REDACTED]							

Investigation Number	Date Issued	Lot Number	Product No.	Product Name	Reason for Investigation	Status	Closed Date
Digitek (digoxin tablets, USP) 0.125 mg							
07-093	12/05/07	70924A1	145	Digitek (digoxin tablets, USP) 0.125 mg	Two tablets of Digoxin tablets 0.125 mg were found with approx. double the thickness from counter channels during packaging/ filling operation.	closed	1/25/2008
08-011	1/29/2008	80044A1	145	Digitek (digoxin tablets, USP) 0.125 mg	One stainless steel screw was found in the tablet well of the BOSS-PACK filling machine during packaging.	closed	3/3/2008
08-017	2/8/2008	80051A	145	Digitek (digoxin tablets, USP) 0.125 mg	During compression of drums #5 and #6 the compression operator observed oil spots on some of the in-process tablets.	closed	3/6/2008
08-028	2/20/2008	80053A	145	Digitek (digoxin tablets, USP) 0.125 mg	Compression operator did not record the metal detector test for the end of the run on 2/15/08.	closed	2/29/2008
08-046	3/13/2008	70078A1	145	Digitek (digoxin tablets, USP) 0.125 mg	Zero stability testing was not conducted when CRT stability study was initiated.	open	
Digitek (digoxin tablets, USP) 0.25 mg							
08-023	2/18/2008	80108A1	146	Digitek (digoxin tablets, USP) 0.25 mg	QA Inspector & labeling operator v. verified an incorrect barcode sticker label.	closed	2/20/2008
08-029	2/20/2008	80152A 80153A	146	Digitek (digoxin tablets, USP) 0.25 mg	Process validation protocols were issued for these batches and were not required. As a result additional samples were taken that were not required.	closed	3/14/2008
08-030	2/20/2008	80133A	146	Digitek (digoxin tablets, USP) 0.25 mg	Operator noticed tablets that were thinner than a typical tablet during the inspection of drum # 2.	closed	3/7/2008
08-031	2/25/2008	80132A 80153A 80154A 80155A	146	Digitek (digoxin tablets, USP) 0.25 mg	The compressed tablet description of MPR # RV-14601 revision 00 is incorrect. The blend description in both the process validation and cleaning protocols is incorrect.	closed	3/13/2008

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